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What is claimed is:

1. A substantially purified human integral membrane protein comprising the amino acid sequence of SEQ ID NO:1 or fragments thereof.

2. A substantially purified variant of human integral membrane protein having at least 90% amino acid identity to SEQ ID NO:1 and which retains at least one functional characteristic of human integral membrane protein.

An isolated and purified polynucleotide sequence encoding the human integral membrane protein of claim 1 or fragments or variants of said polynucleotide sequence.

- 4. A composition comprising the polynucleotide sequence of claim 3.
- 5. A polynucleotide sequence which hybridizes to the polynucleotide sequence of claim 3.
- 6. A polynucleotide sequence which is complementary to the polynucleotide sequence of claim 3 or fragments or variants thereof.
- 7. An isolated and purified polynucleotide sequence comprising SEQ ID NO:2 or fragments or variants thereof.
- 8. A polynucleotide sequence which is complementary to the polynucleotide sequence of claim 7.
 - 9. An expression vector containing at least a fragment of the polynucleotide sequence of claim 3.
- 30 10. A host cell containing the vector of claim 9.

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- 11. A method for producing a polypeptide comprising the amino acid sequence of SEQ ID NO:1, or a fragment thereof, the method comprising the steps of:
 - a) culturing the host cell of claim 10 under conditions suitable for the expression of the polypept de, and
 - b) recovering the polypeptide from the host cell culture.
- 12. A pharmac eutical composition comprising a substantially purified human integral membrane protein having the amino acid sequence of SEQ ID NO:1 in conjunction with a suitable pharmac eutical carrier.
 - 13. A purified antibody which specifically binds to the polypeptide of claim 1.
 - 14. A purified agonist of the polypeptide of claim 1.
 - 15. A purified antagonist of the polypeptide of claim 1.
- 16. A method for treating cancer comprising administering to a subject in need of such treatment an effective amount of the pharmaceutical composition of claim 12.
- 17. A method for treating a neuronal disorder comprising administering to a subject in need of such treatment an effective amount of the antagonist of claim 15.
- 18. A method for treating an immunological disorder comprising administering to a subject in need of such treatment an effective amount of the antagonist of claim 15
- 19. A method for detecting a polynucleotide which encodes human integral membrane protein in a biological sample comprising the steps of:
 - a) hybridizing the polynucleotide of claim 6 to nucleic acid material of a biological sample, thereby forming a hybridization complex; and
 - b) detecting said hybridization complex, wherein the presence of said

PF-0339 US

complex correlates with the presence of a polynucleotide encoding human integral membrane protein in said biological sample.

The method of claim 10 wherein the nucleic acid material is amplified by the polymerase chain reaction prior to hybridization.

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